FDA 510(k) Premarket Notification

K875038 MENTOR UNISTENT URETERAL STENT SET

CLEARED

Mar 3, 1988
87 days to decision

K875038 · Product code: **FAD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k875038/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Dec 7, 1987
Decision date	Mar 3, 1988
Days to decision	87 days
Third-party review	No

APPLICANT

Company	**Mentor Corp.**
Location	Mchenry, IL, US
Contact	R BRECKENRIDGE
510(k) history	61 submissions · 61 cleared · 1977-2013

Mentor Corp. is a surgical aesthetics and medical device company based in McHenry, US. Now part of Johnson & Johnson MedTech, the brand supplies products to plastic surgeons and specialists worldwide. Mentor has received FDA 510(k) clearances from total submissions since its first clearance in 1977. The company's regulatory record spans General & Plastic Surgery, Gastroenterology & Urology, Obstetrics & Gynecology, and Radiology device categories. The latest clearance was recorded in 2013, reflecting the company's historical significance in surgical device innovation. Men...